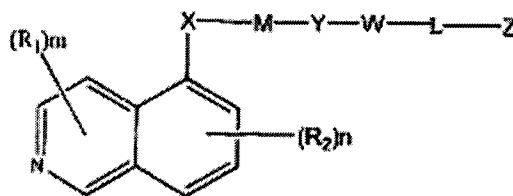


## Amendments to the Claims

The following listing of claims replaces all prior listings and version of claims in this application.

1. (Previously Presented) A compound of Formula I:



Formula I

wherein:

$R_1$  and  $R_2$  are independently selected from the group consisting of hydrogen, a lower alkyl group, a lower alkoxy group, substituted or unsubstituted phenyl group, a lower alkyl substituted with at least one substituent selected from the group consisting of a phenyl group, a halogen, hydroxyl, thiol, nitro, cyano, or amino group;  $m$  and  $n$  are each independently 0-3;

$X$  is selected from the group consisting of  $SO_2-NH$ ,  $S$  and  $O$ ;

$M$  represents substituted or unsubstituted alkylene of 1-4 carbon atoms;

$Y$  is selected from the group consisting of amide, amine, urea, carbamate, hydrazine or sulfonamide;

$Z$  is  $Arg-Pro-Arg-R_4-R_5-R_6-R_7$ ;

$R_4$ ,  $R_5$ , and  $R_6$  are each independently selected from the group consisting of threonine, serine, glutamic acid allyl ester, homocitrulline, lysine, methionine, norleucine, ornithine, arginine, glycine, diaminopropionic acid, diaminobutyric acid,  $GlyNH_2$ , and alanine; or are an  $N^\alpha$ - $\omega$ -functionalized derivative of an amino acid selected from the group of glycine, alanine and tyrosine;

$R_7$  is selected from the group consisting of phenylalanine, homoleucine, norleucine, glutamic acid allyl ester;

W may be absent so that Y is connected to L or R<sub>4</sub>, or W is N-(8-sulfonamide-5-isoquinoline) ethylenediamine; and

L may be absent so that W (if present) or Y is connected to R<sub>4</sub>, or L is selected from the group consisting of glycine,  $\beta$ -alanine, phenylalanine, aminobutyric acid and aminopentanoic acid and connects W (if present) or Y with R<sub>4</sub>.

2. (Previously Presented) The compound of claim 1 wherein, in Formula I:

R<sub>1</sub> and R<sub>2</sub> are independently selected from the group consisting of methyl, ethyl, ethoxy and dimethylamine;

m and n are each 1;

M represents substituted or unsubstituted alkylene of 2 carbon atoms; and

Y is selected from the group consisting of amide and amine.

Claims 3 to 6. (Cancelled)

7. (Previously Presented) The compound according to claim 1 wherein the compound comprises:

Arg-Pro-Arg-Thr-Glu-(bAla-5-mercaptoaminopropyl-isoquinoline)-Ser-Phe (SEQ ID NO: 3).

8. (Previously Presented) The compound according to claim 1 wherein the compound comprises:

Arg-Pro-Arg-Thr-Glu-(5-mercaptoaminopropyl-isoquinoline)-Ser-Phe (SEQ ID NO: 4).

9. (Previously Presented) The compound according to claim 1 wherein the compound comprises:

Arg-Pro-Arg-Orn-Glu-(5-aminoethylsulfonamide-isoquinoline)-Ser-Phe (SEQ ID NO: 5).

10. (Previously Presented) The compound according to claim 1 wherein the compound comprises:

Arg-Pro-Arg-Nva-Glu-(5-mercaptoaminopropyl-isoquinoline)-Ser-Phe (SEQ ID NO: 6).

11. (Previously Presented) The compound according to claim 1 wherein the compound comprises:

Arg-Pro-Arg-Nle-Glu-(5-mercaptoaminopropyl-isoquinoline)-Ser-Phe (SEQ ID NO: 7).

12. (Previously Presented) The compound according to claim 1 wherein the compound comprises:

Arg-Pro-Arg-Orn-Glu-(Gly-5-aminoethylsulfonamide)-Dab-Hol (SEQ ID NO: 8).

13. (Previously Presented) The compound according to claim 1 wherein the compound comprises:

Arg-Pro-Arg-Nle-Glu-(Gly-5-aminoethylsulfonamide)-Dab-Phe (SEQ ID NO: 9).

14. (Currently Amended) The compound according to claim 1 wherein the compound [~~comprise~~] comprises:

Arg-Pro-Arg-Nle-Glu-(Gly-5-aminoethylsulfonamide)-Dab-Hol (SEQ ID NO: 10).

15. (Original) A pharmaceutical composition comprising as an active ingredient a compound according to claim 1, and a pharmaceutically acceptable diluent or carrier.

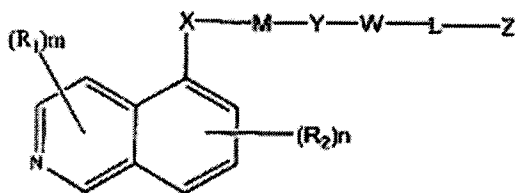
16. (Original) A protein kinase inhibitor comprising as an active ingredient a compound according to claim 1, and a pharmaceutically acceptable diluent or carrier.

17. (Previously Presented) A method of treatment of diabetes, hemorrhagic shock, or inflammatory disease, comprising administering to a patient in need thereof a therapeutically effective amount of a compound according to claim 1.

18. (Cancelled)

19. (Cancelled)

20. (Currently Amended) A compound of Formula I:



Formula I

wherein:

~~[[R1]]~~  $R_1$  and ~~[[R2]]~~  $R_2$  are independently selected from the group consisting of hydrogen, a lower alkyl group, a lower alkoxy group, substituted or unsubstituted phenyl group, a lower alkyl substituted with at least one substituent selected from the group consisting of a phenyl group, a halogen, hydroxyl, thiol, nitro, cyano, or amino group; m and n are each independently 0-3;

X is selected from the group consisting of ~~[[S02]]~~  $SO_2$ -NH, S and O;

M represents substituted or unsubstituted alkylene of 1-4 carbon atoms;

Y is selected from the group consisting of amide, amine, urea, carbamate, hydrazine or sulfonamide;

W is absent or is selected from the group consisting of substituted or unsubstituted alkylene, aliphatic, aromatic or heterocyclic moiety, of 1-18 carbon atoms;

L is absent or is selected from the group consisting of amide, amine, urea, carbamate, hydrazine or sulfonamide; and

Z is a peptide or peptidomimetic moiety comprising one of the following sequences:

Arg-Pro-Arg-Thr-Glu-Ser-Phe (SEQ ID NO: 3);

Arg-Pro-Arg-Thr-Glu-Ser-Phe (SEQ ID NO: 4);

Arg-Pro-Arg-~~[[Om]]~~  $Orn$ -Glu-Ser-Phe (SEQ ID NO: 5);

Arg-Pro-Arg-Nva-Glu-Ser-Phe (SEQ ID NO: 6);

Arg-Pro-Arg-Nle-Glu-Ser-Phe (SEQ ID NO: 7);

Arg-Pro-Arg-~~[[Om]]~~  $Orn$ -Glu-Dab-Hol (SEQ ID NO: 8);

Arg-Pro-Arg-Nle-Glu-Dab-Phe (SEQ ID NO: 9); or

Arg-Pro-Arg-Nle-Glu-Dab-Hol (SEQ ID NO: 10); and wherein Y, W if present, or L if present are linked to the Glu residues of the sequences.

21. (Previously Presented) A pharmaceutical composition comprising as an active ingredient a compound according to claim 20, and a pharmaceutically acceptable diluent or carrier.

22. (Previously Presented) A protein kinase inhibitor comprising as an active ingredient a compound according to claim 20, and a pharmaceutically acceptable diluent or carrier.

23. (Previously Presented) A method of treatment of diabetes, hemorrhagic shock, or inflammatory disease, comprising administering to a patient in need thereof a therapeutically effective amount of a compound according to claim 20.

24. (Cancelled)